

LISTING OF THE CLAIMS

This listing of claims replaces all prior versions, and listings of claims in the application:

1. (Withdrawn) A material that can be reconstituted to provide a replacement fluid for use in treating a joint malady of an animal made by the process comprising:
 - collecting synovial from donor animals;
 - removing impurities, cellular and pathogenic components from said synovial fluid to create a purified synovial fluid; and,
 - lyophilizing said purified synovial fluid.
2. (Withdrawn) The material of claim 1 wherein said process of collecting synovial from donor animals further comprises:
 - selecting a donor joint from said donor animal;
 - injecting a joint capsule of said selected donor joint with and needle attached to a syringe;
 - aspirating fluid joint contents into said syringe; and,
 - preserving said fluid joint contents for further processing.
3. (Withdrawn) The material of claim 2 wherein said process of preserving said fluid joint contents for further processing further comprises:
 - freezing said fluid joint contents at a temperature less than zero degrees Centigrade.
4. (Withdrawn) The material of claim 1 wherein said process of removing impurities, cellular and pathogenic components from said synovial fluid to create a purified synovial fluid further comprises:
 - separating higher density particles within said synovial fluid by centrifuge;
 - removing said higher density particles from a supernate of said synovial fluid; and,
 - filtering said supernate to remove additional particulates greater than approximately .45 mm in size.

5. (Withdrawn) The material of claim 1 wherein said process of lyophilizing said synovial fluid further comprises:
 - stabilizing said purified synovial fluid;
 - freezing said purified synovial fluid at approximately -45 degrees Centigrade
 - reducing the ambient air pressure to said purified synovial fluid to less than 50 microns of mercury;
 - maintaining said purified synovial fluid at -35 degrees Centigrade for approximately 72 hours;
 - maintaining said purified synovial fluid at 0 degrees Centigrade for approximately 12 hours; and,
 - maintaining said purified synovial fluid at 25 degrees Centigrade for approximately 12 hours.
6. (Withdrawn) The material of claim 5 wherein said process of lyophilizing said synovial fluid further comprises:
 - placing said purified synovial fluid under vacuum.
7. (Withdrawn) The material of claim 1 further comprising the process of:
 - providing said lyophilized synovial fluid to users for reconstitution as an intraarticular injection in an aseptic manner.
8. (Withdrawn) The material of claim 7 wherein said process of providing said lyophilized synovial fluid to users for reconstitution as an intraarticular injection in an aseptic manner further comprises:
 - providing said lyophilized synovial fluid in a vacuum-sealed vial for reconstitution within said vial to produce a single-use intraarticular injection.
9. (Currently amended) A method of treating a joint malady of an animal comprising:
 - intraarticularly injecting a replacement fluid in the joint space of said animal, said replacement fluid comprising synovial fluid that has been harvested from other animals ~~that and~~ has been processed lyophilized, packaged and reconstituted.

10. (Original) A method of treating a joint malady of an animal by intraarticularly injecting a purified synovial fluid in the joint space of said animal, said purified synovial fluid made by the process of:
 - collecting synovial from donor animals;
 - removing impurities, cellular and pathogenic components from said synovial fluid to create a purified synovial fluid;
 - lyophilizing said purified synovial fluid; and,
 - reconstituting said purified synovial fluid to approximately its original volume.
11. (Original) A method of claim 10 wherein said step of collecting synovial from donor animals further comprises:
 - selecting a donor joint from said donor animal;
 - injecting a joint capsule of said selected donor joint with and needle attached to a syringe;
 - aspirating fluid joint contents into said syringe; and,
 - preserving said fluid joint contents for further processing.
12. (Currently amended) A method of claim 11 wherein said step of preserving said fluid joint contents for further processing further comprises:
 - freezing said fluid joint contents at temperature less ~~that~~ than zero degrees Centigrade.
13. (Original) A method of claim 10 wherein said step of removing impurities, cellular and pathogenic components from said synovial fluid to create a purified synovial fluid further comprises:
 - separating higher density particles within said synovial fluid by centrifuge;
 - removing said higher density particles from a supernate of said synovial fluid;
 - and,
 - filtering said supernate to remove additional particulates greater than approximately .45 mm in size.

14. (Original) A method of claim 10 wherein said step of lyophilizing said synovial fluid further comprises:
- stabilizing said purified synovial fluid;
 - freezing said purified synovial fluid at approximately -45 degrees Centigrade
 - reducing the ambient air pressure to said purified synovial fluid to less than 50 microns of mercury;
 - maintaining said purified synovial fluid at -35 degrees Centigrade for approximately 72 hours;
 - maintaining said purified synovial fluid at 0 degrees Centigrade for approximately 12 hours; and,
 - maintaining said purified synovial fluid at 25 degrees Centigrade for approximately 12 hours.
15. (Original) A method of claim 14 wherein said step of lyophilizing said synovial fluid further comprises:
- placing said purified synovial fluid under vacuum.
16. (Currently amended) A method of claim 10 further comprising the step of:
- providing said lyophilized synovial fluid to users for reconstitution as an intraarticular injection in an aseptic manner;
17. (Original) A method of claim 16 wherein said step of providing said lyophilized synovial fluid to users for reconstitution as an intraarticular injection in an aseptic manner further comprises:
- providing said lyophilized synovial fluid in a vacuum-sealed vial for reconstitution within said vial to produce a single-use intraarticular injection.
18. (Withdrawn) A method of manufacturing a concentrated material that can be reconstituted to provide a replacement fluid for use in treating a joint malady of an animal comprising:
- collecting synovial from donor animals,

removing impurities, cellular and pathogenic components from said synovial fluid;

lyophilizing said purified synovial fluid to form a dense concentrate; and,

packaging said lyophilized concentrate in a manner so as to provide said lyophilized concentrate in a form that is reconstitutable to serve as an injectable replacement fluid.

19. (Withdrawn) A method of claim 18 wherein said step of collecting synovial from donor animals further comprises:

selecting a donor joint from said donor animal;

injecting a joint capsule of said selected donor joint with and needle attached to a syringe;

aspirating fluid joint contents into said syringe; and,

preserving said fluid joint contents for further processing.

20. (Withdrawn) A method of claim 19 wherein said step of preserving said fluid joint contents for further processing further comprises:

freezing said fluid joint contents at temperature less than zero degrees Centigrade.

21. (Withdrawn) A method of claim 18 wherein said step of removing impurities, cellular and pathogenic components from said synovial fluid to create a purified synovial fluid further comprises:

separating higher density particles within said synovial fluid by centrifuge;

removing said higher density particles from a supernate of said synovial fluid;
and,

filtering said supernate to remove additional particulates greater than approximately .45 mm in size.

22. (Withdrawn) A method of claim 18 wherein said step of lyophilizing said synovial fluid further comprises:

stabilizing said purified synovial fluid;

freezing said purified synovial fluid at approximately -45 degrees Centigrade
reducing the ambient air pressure to said purified synovial fluid to less than 50
microns of mercury;

maintaining said purified synovial fluid at -35 degrees Centigrade for
approximately 72 hours;

maintaining said purified synovial fluid at 0 degrees Centigrade for approximately
12 hours; and,

maintaining said purified synovial fluid at 25 degrees Centigrade for
approximately 12 hours.

23. (Withdrawn) A method of claim 22 wherein said step of lyophilizing said synovial fluid
further comprises:

placing said purified synovial fluid under vacuum.

24. (Withdrawn) A method of claim 18 further comprising the step of:

providing said lyophilized synovial fluid to users for reconstitution as an
intraarticular injection in an aseptic manner:

25. (Withdrawn) A method of claim 24 wherein said step of providing said lyophilized
synovial fluid to users for reconstitution as an intraarticular injection in an aseptic manner
further comprises:

providing said lyophilized synovial fluid in a vacuum-sealed vial for
reconstitution within said vial to produce a single-use intraarticular injection.